



US009087687B2

(12) **United States Patent**
Adam et al.

(10) **Patent No.:** **US 9,087,687 B2**
(45) **Date of Patent:** **Jul. 21, 2015**

(54) **THIN HETEROSTRUCTURE CHANNEL DEVICE**

USPC 257/192, 369, E21.409, E21.632,
257/E27.062; 438/218
See application file for complete search history.

(75) Inventors: **Thomas N. Adam**, Slingerlands, NY (US); **Kangguo Cheng**, Guilderland, NY (US); **Hong He**, Schenectady, NY (US); **Ali Khakifirooz**, Slingerlands, NY (US); **Alexander Reznicek**, Mount Kisco, NY (US)

(56) **References Cited**

(73) Assignee: **INTERNATIONAL BUSINESS MACHINES CORPORATION**, Armonk, NY (US)

U.S. PATENT DOCUMENTS

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 601 days.

5,426,069	A	6/1995	Selvakumar et al.	
5,461,250	A	10/1995	Burghartz et al.	
6,713,779	B2	3/2004	Tezuka et al.	
7,648,868	B2	1/2010	Majumdar et al.	
8,008,138	B2*	8/2011	Cheng et al.	438/157
2004/0136866	A1*	7/2004	Pontis et al.	422/57
2004/0197975	A1	10/2004	Krivokapic et al.	
2005/0148161	A1*	7/2005	Chen et al.	438/478

(Continued)

(21) Appl. No.: **13/336,251**

OTHER PUBLICATIONS

(22) Filed: **Dec. 23, 2011**

K. Cheng et al., "Fully depleted extremely thin SOI technology fabricated by a novel integration scheme featuring implant-free, zero-silicon-loss, and faceted raised source/drain," 2009 Symposium on VLSI Technology, Jun. 16-18, 2009, pp. 212-213.

(65) **Prior Publication Data**

(Continued)

US 2013/0161693 A1 Jun. 27, 2013

(51) **Int. Cl.**

Primary Examiner — Vongsavanh Sengdara

H01L 21/8238 (2006.01)

(74) *Attorney, Agent, or Firm* — Scully, Scott, Murphy & Presser, P.C.; Daniel P. Morris, Esq.

H01L 21/02 (2006.01)

H01L 21/84 (2006.01)

H01L 27/12 (2006.01)

H01L 21/3065 (2006.01)

H01L 29/10 (2006.01)

(52) **U.S. Cl.**

(57) **ABSTRACT**

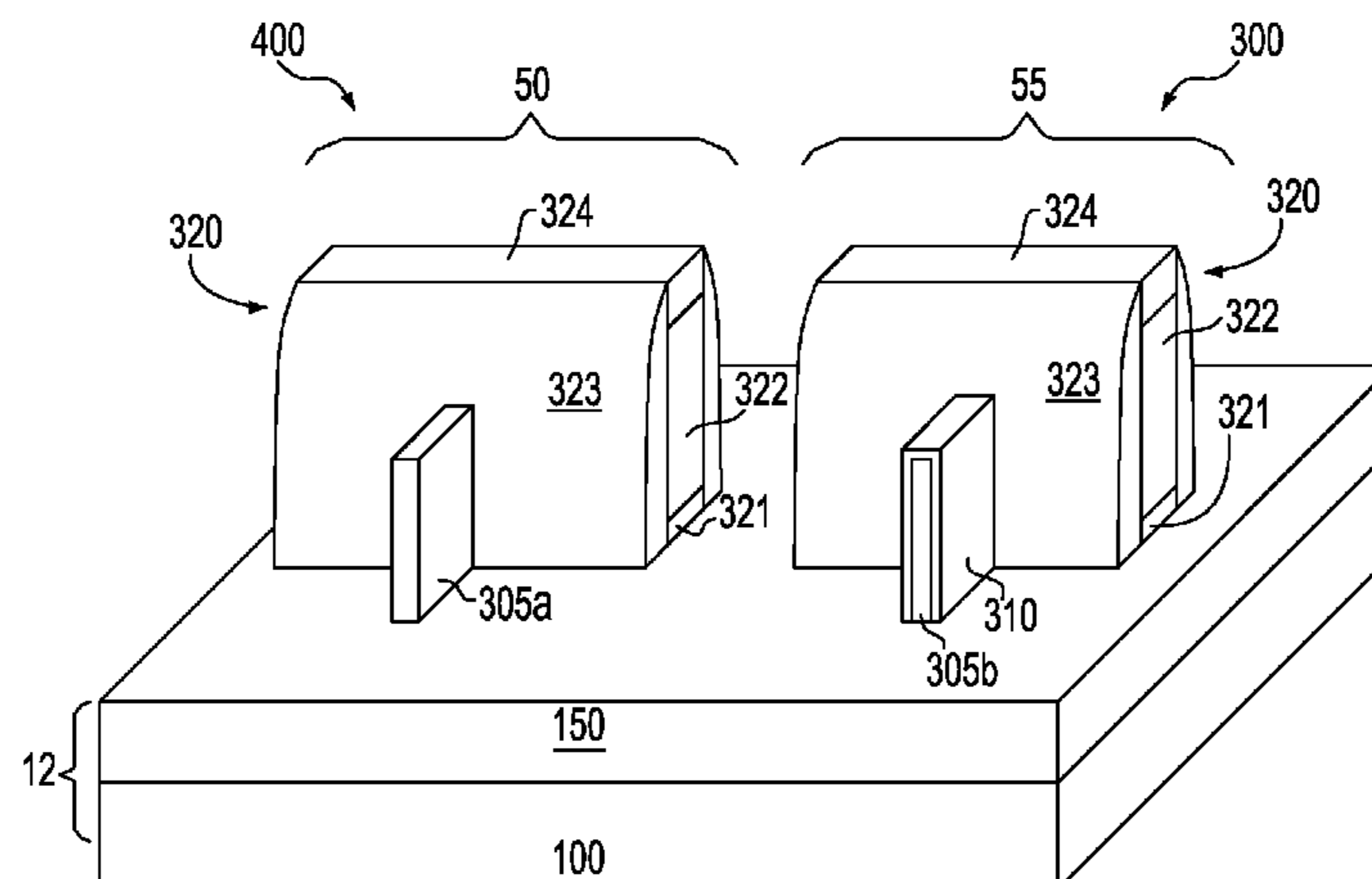
CPC **H01L 21/02057** (2013.01); **H01L 21/3065** (2013.01); **H01L 21/823807** (2013.01); **H01L 21/823821** (2013.01); **H01L 21/845** (2013.01); **H01L 27/1211** (2013.01); **H01L 29/1054** (2013.01); **H01L 21/0262** (2013.01); **H01L 21/02532** (2013.01)

A method of fabricating a semiconductor device that includes providing a substrate having at least a first semiconductor layer atop a dielectric layer, wherein the first semiconductor layer has a first thickness of less than 10 nm. The first semiconductor layer is etched with a halide based gas at a temperature of less than 675° C. to a second thickness that is less than the first thickness. A second semiconductor layer is epitaxially formed on an etched surface of the first semiconductor layer. A gate structure is formed directly on the second semiconductor layer. A source region and a drain region is formed on opposing sides of the gate structure.

(58) **Field of Classification Search**

CPC H01L 21/823807; H01L 21/823821; H01L 21/845; H01L 27/1211; H01L 29/1054

11 Claims, 4 Drawing Sheets



(56)

References Cited

U.S. PATENT DOCUMENTS

2006/0166416	A1 *	7/2006	Dalton et al.	438/153
2008/0054364	A1 *	3/2008	Hokazono	257/369
2008/0135886	A1	6/2008	Irisawa et al.	
2008/0191286	A1	8/2008	Chang et al.	
2010/0184260	A1	7/2010	Luo et al.	
2010/0200937	A1	8/2010	Bedell et al.	
2010/0203737	A1 *	8/2010	Morikawa et al.	438/719
2011/0042744	A1	2/2011	Cheng et al.	
2011/0086475	A1	4/2011	Yamazaki et al.	
2011/0147840	A1	6/2011	Cea et al.	
2011/0159668	A1 *	6/2011	Dhumal et al.	438/479
2011/0227157	A1 *	9/2011	Yang et al.	257/347
2011/0248348	A1	10/2011	Gan et al.	
2012/0074386	A1	3/2012	Rackmady et al.	
2012/0091538	A1	4/2012	Lin et al.	
2012/0261754	A1 *	10/2012	Cheng et al.	257/347

OTHER PUBLICATIONS

K. Cheng et al., "Extremely thin SOI (ETSOI) CMOS with record low variability for low power system-on-chip applications," 2009 IEEE International Electron Devices Meeting (IEDM), Dec. 7-9, 2009, 4 pages.

M. Kim et al., Increased critical thickness for high Ge-content strained SiGe-on-Si using selective epitaxial growth, Appl. Phys. Lett., vol. 97, Issue 26, 2010, pp. 262106-1-262106-3. Dec. 27, 2010.

S. Baudot et al., "Fully Depleted Strained Silicon-on-Insulator p-MOSFETs With Recessed and Embedded Silicon-Germanium Source/Drain," IEEE Electron Device Letters, vol. 31, Issue:10, Oct. 2010, Date of Publication: Aug. 23, 2010, pp. 1074-1076.

International Search Report and Written Opinion dated Feb. 7, 2013 issued in PCT/US2012/66928.

Office Action dated Jul. 28, 2014 received in U.S. Appl. No. 13/607,875.

Final Office Action dated Jan. 29, 2015 received in U.S. Appl. No. 13/607,875.

* cited by examiner

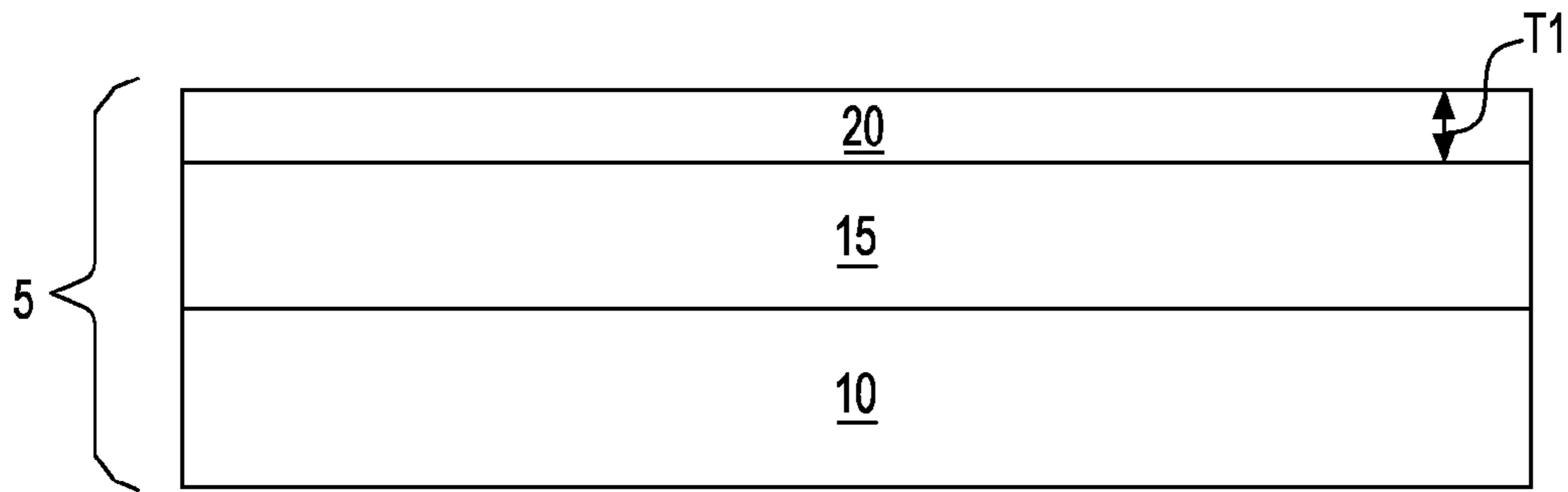


FIG. 1

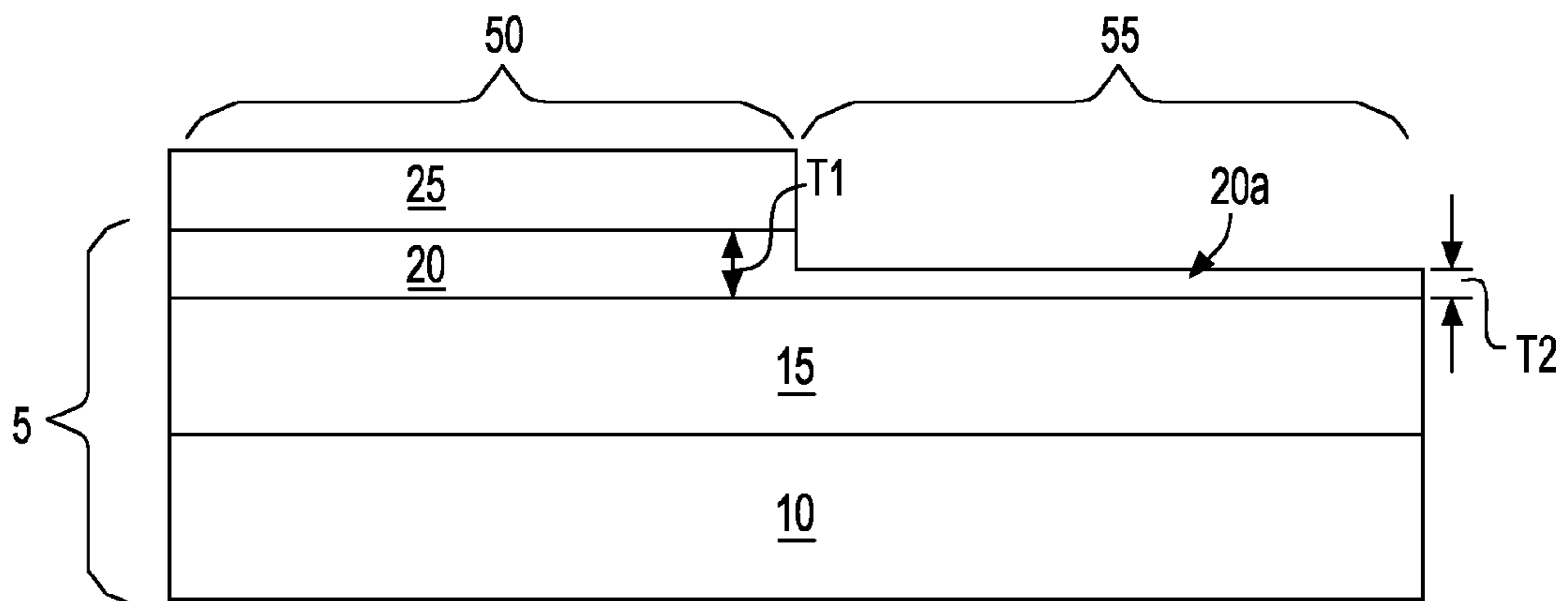


FIG. 2

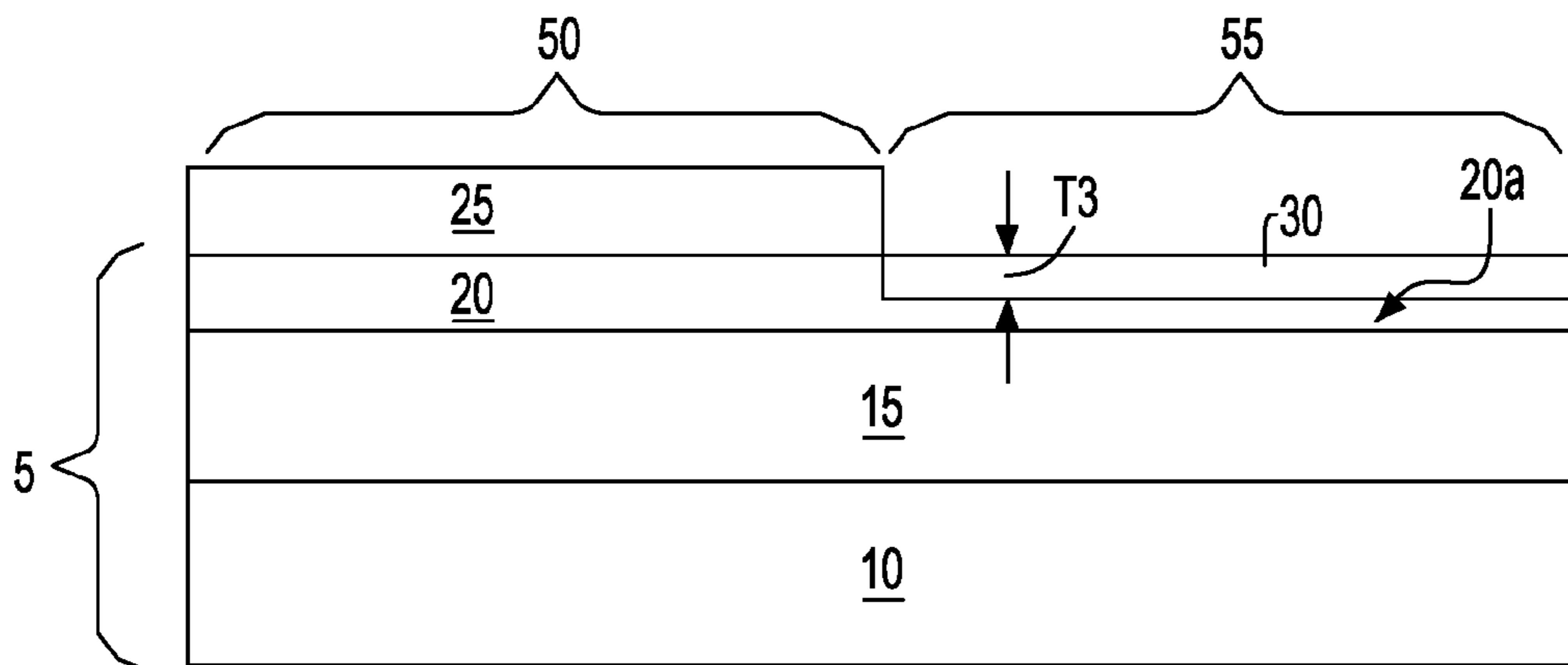


FIG. 3

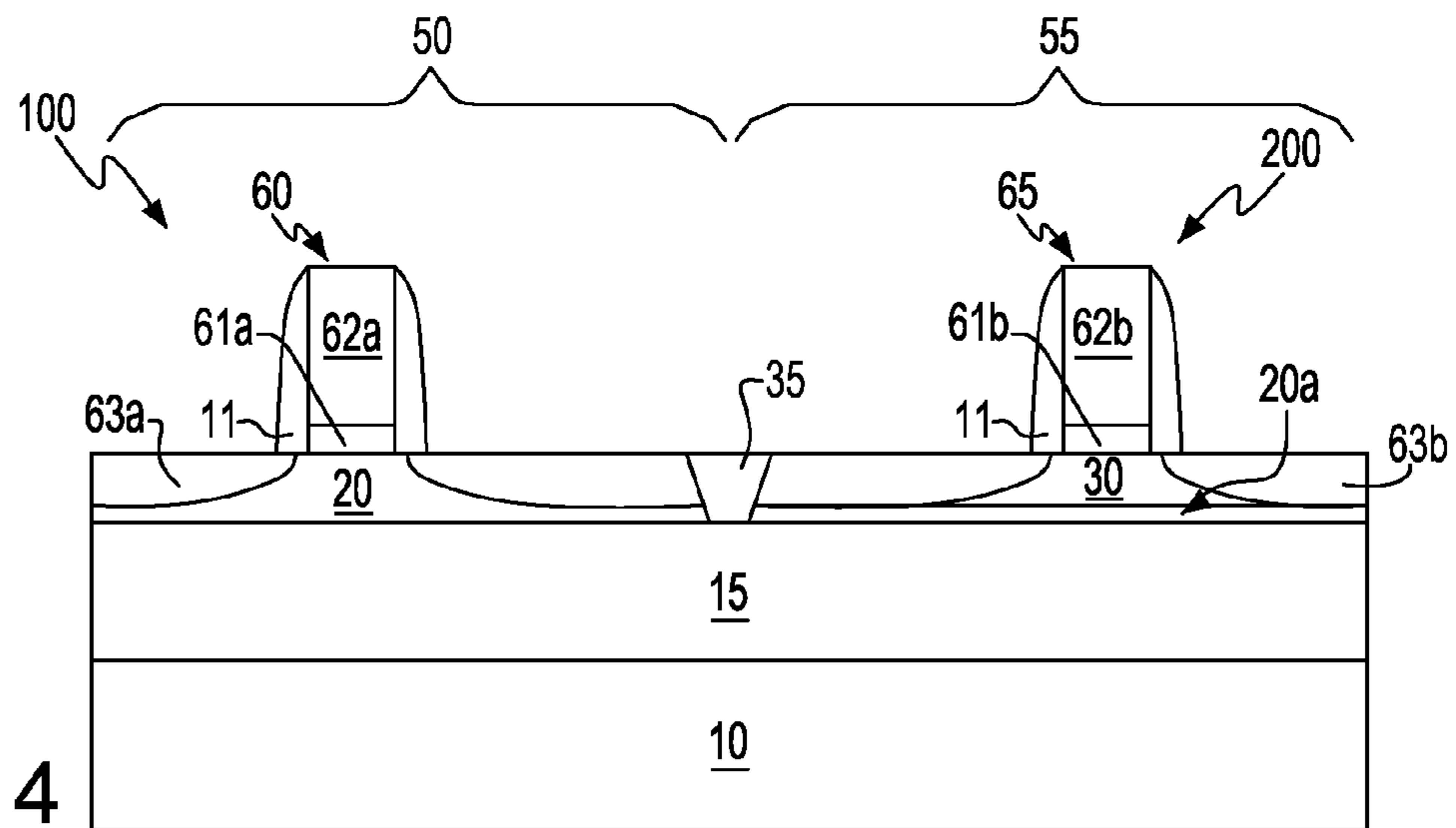


FIG. 4

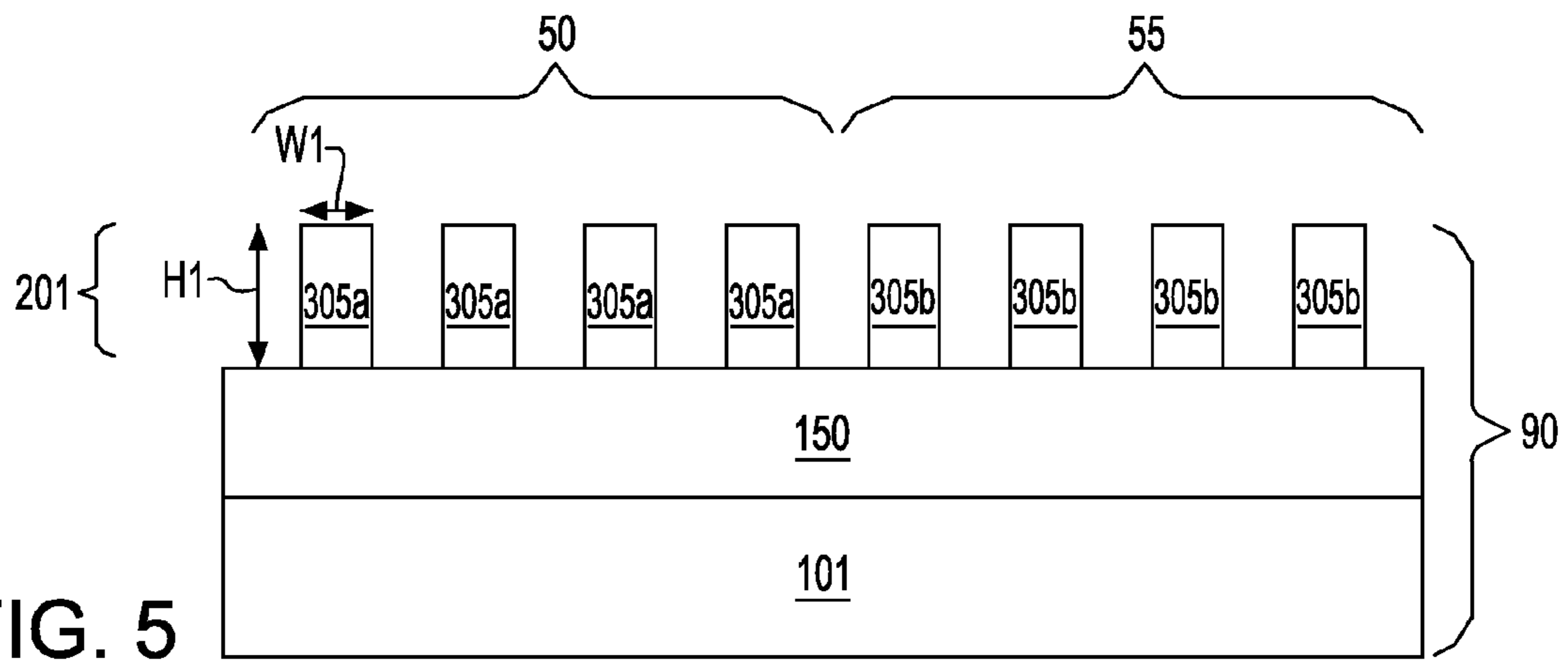


FIG. 5

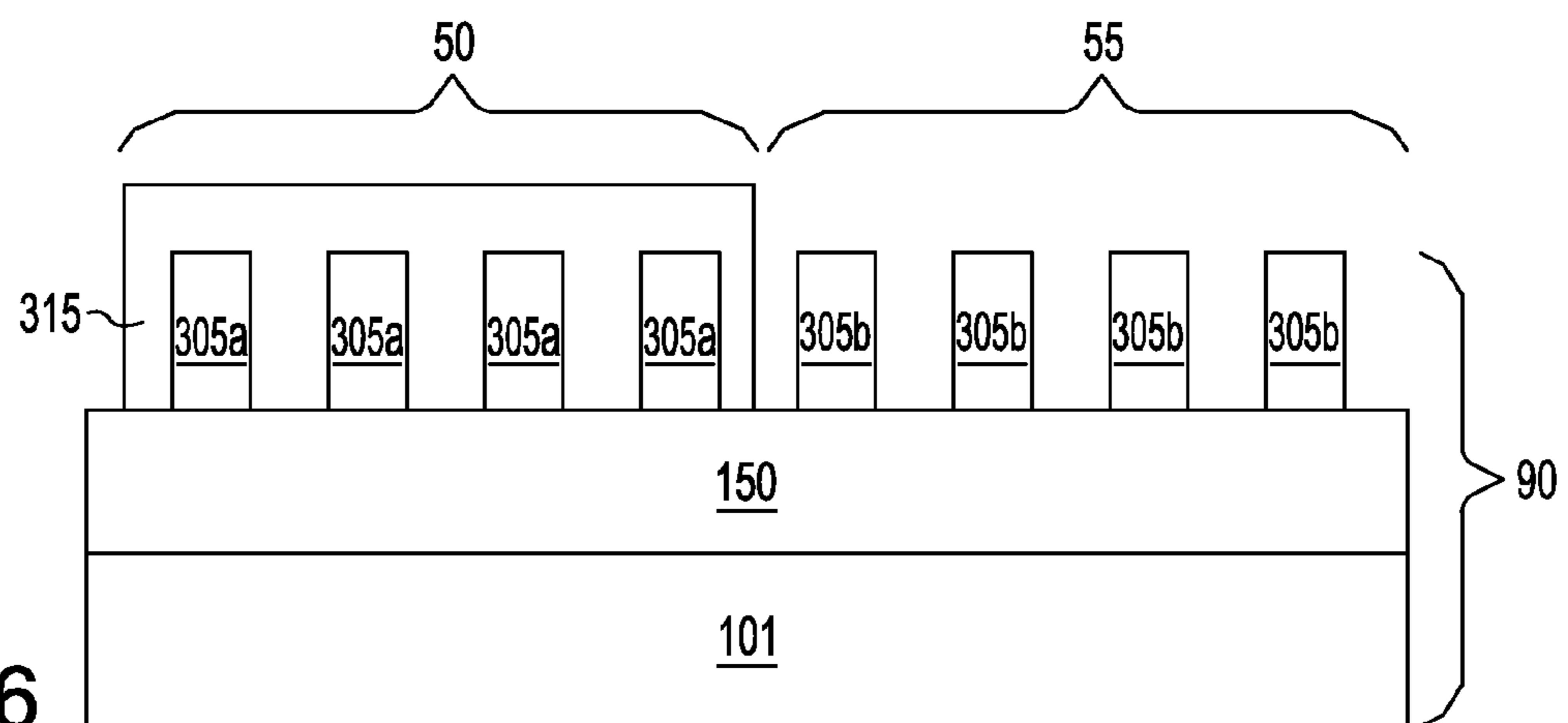


FIG. 6

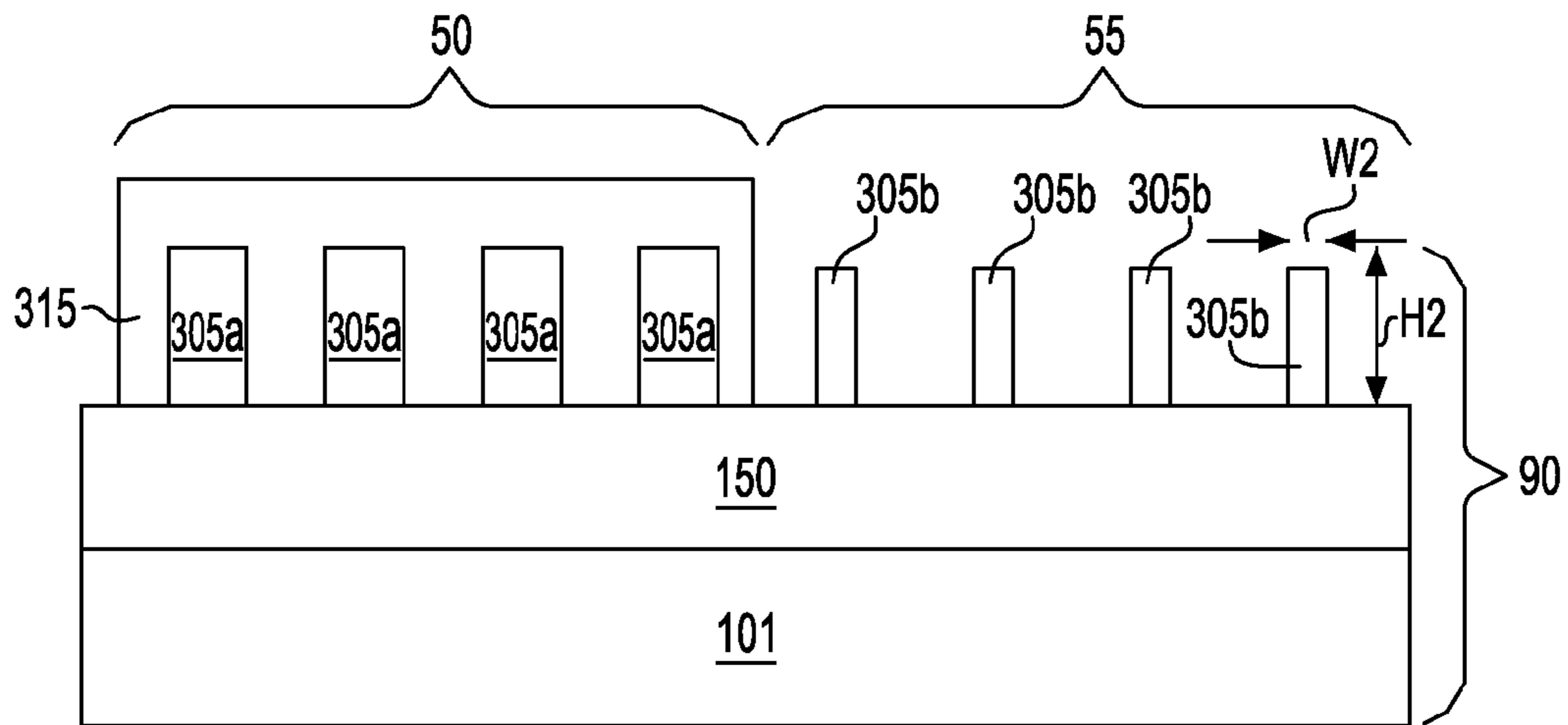


FIG. 7

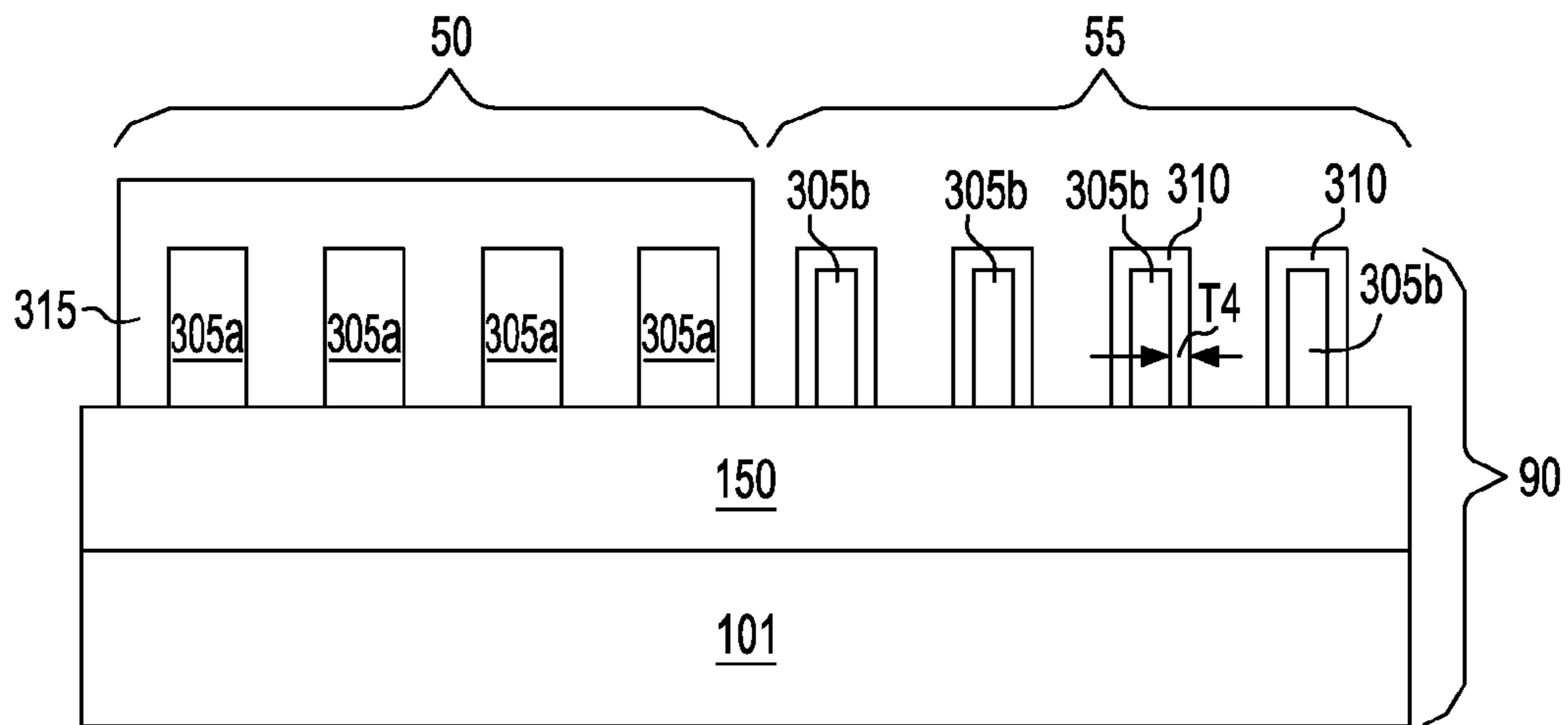


FIG. 8

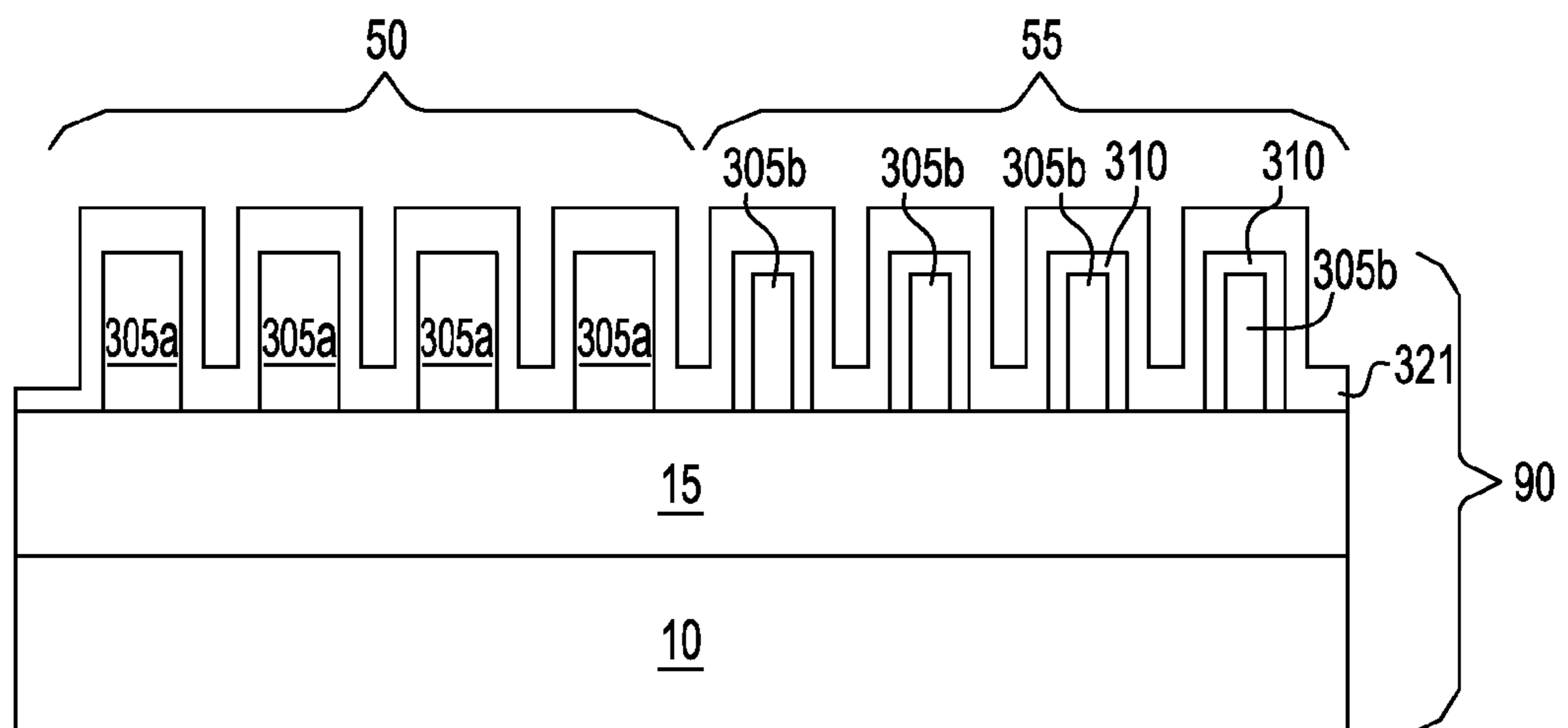


FIG. 9

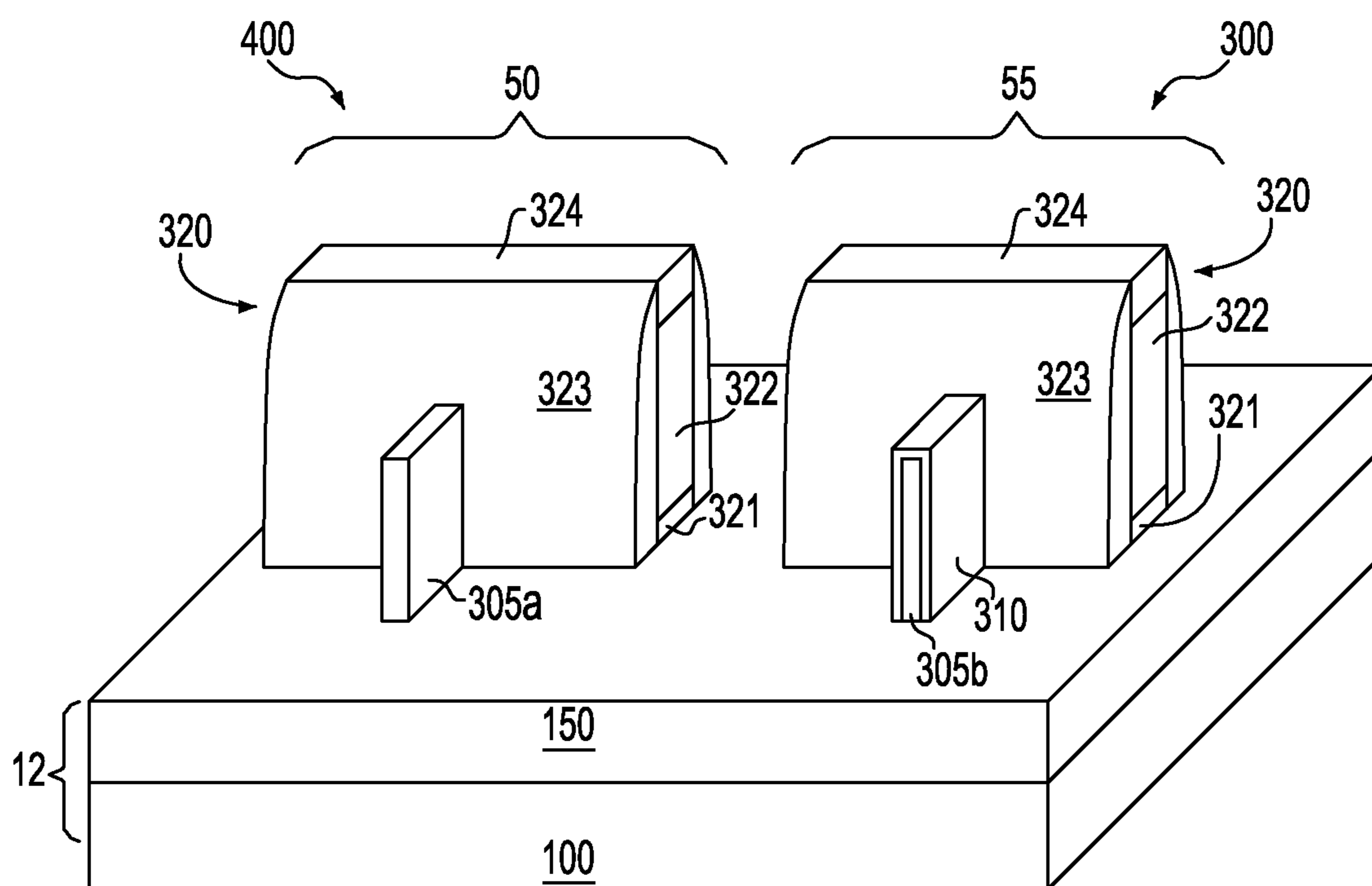


FIG. 10

THIN HETEROSTRUCTURE CHANNEL DEVICE

BACKGROUND

The present disclosure relates generally to semiconductor devices. More particularly, the present disclosure relates to scaling of semiconductor devices.

In order to be able to make integrated circuits (ICs), such as memory, logic, and other devices, of higher integration density than currently feasible, one has to find ways to further downscale the dimensions of field effect transistors (FETs), such as metal-oxide-semiconductor field effect transistors (MOSFETs) and complementary metal oxide semiconductors (CMOS). Scaling achieves compactness and improves operating performance in devices by shrinking the overall dimensions and operating voltages of the device while maintaining the device's electrical properties. Typically, all dimensions of the device must be scaled simultaneously in order to optimize the electrical performance of the device.

SUMMARY

A method of fabricating a semiconductor device is provided that, in one embodiment, may begin with providing a substrate comprising at least a first semiconductor layer atop a dielectric layer, in which the first semiconductor layer is comprised of a first semiconductor material and has a first thickness of less than 10 nm. The first semiconductor layer may be etched with a halide based gas at a temperature of less than 675° C. to a second thickness that is less than the first thickness. A second semiconductor layer may be epitaxially formed on the etched surface of the first semiconductor layer. A gate structure can be formed directly on the second semiconductor layer, and source regions and drain regions may be formed on opposing sides of the gate structure.

In another aspect of the present disclosure, a complementary metal oxide semiconductor (CMOS) device is provided that includes a substrate having a first device region and a second device region. In one embodiment, the first device region includes a first thickness silicon layer that is present on a first portion of a buried insulating layer. In one embodiment, the second device region includes a silicon germanium layer on a second thickness silicon layer, wherein the second thickness silicon layer is present on a second portion of the buried insulating layer. The second thickness silicon layer has a lesser thickness than the first thickness silicon layer. A planar n-type semiconductor device may be present in the first device region of the substrate, and a planar p-type semiconductor device may be present in the second device region of the substrate.

In yet another aspect of the present disclosure, a method of forming a fin field effect transistor (finFET) is provided. In one embodiment, the method of forming the finFET includes providing at least one fin structure having a width of less than 20 nm, wherein the at least one fin structure is composed of a first semiconductor material. The at least one fin structure is then etched with a halide based gas at a temperature of less than 675° C. to a second width that is less than the first width of the at least one fin structure. A second semiconductor material is epitaxially formed on an etched surface of the at least one fin structure. A gate structure is formed atop a portion of the second semiconductor material. Source and drain regions may be formed on the exposed sidewalls of the second semiconductor material that is present on the fin structure on each side of the gate structure.

In a further aspect of the present disclosure, a fin field effect transistor (finFET) is provided that includes a fin structure of a first semiconductor material having a width ranging from 1.0 nm to 5.0 nm. A second semiconductor material is present on an upper surface and sidewalls of the fin structure. The second semiconductor material is a conformal epitaxial material having a thickness ranging from 1 nm to 10 nm. A gate structure is present on a portion of the second semiconductor material that is overlying the fin structure, and source and drain regions are present on opposing sides of the gate structure.

DESCRIPTION OF SEVERAL VIEWS OF THE DRAWINGS

The following detailed description, given by way of example and not intended to limit the present disclosure solely thereto, will best be appreciated in conjunction with the accompanying drawings, wherein like reference numerals denote like elements and parts, in which:

FIG. 1 is a side cross-sectional view depicting providing a substrate comprising at least a first semiconductor layer atop a dielectric layer, wherein the first semiconductor layer has a thickness of less than 10 nm, as used in one embodiment of a method for forming a planar semiconductor device, in accordance with the present disclosure.

FIG. 2 is a side cross-sectional view depicting forming an etch mask over a first device region of the substrate and etching an exposed portion of the first semiconductor layer in a second device region, in accordance with one embodiment of the present disclosure.

FIG. 3 is a side cross-sectional view depicting epitaxially forming a second semiconductor layer on the etched surface of the first semiconductor layer, in accordance with one embodiment of the present disclosure.

FIG. 4 is a side cross-sectional view depicting forming a gate structure in each of the first device region and the second device region, and forming source and drain regions on opposing sides of the gate structures, in accordance with one embodiment of the present disclosure.

FIG. 5 is a side cross-sectional view depicting forming fin structures having a width of less than 20 nm, as used in one embodiment of a method for forming a finFETs device, in accordance with the present disclosure.

FIG. 6 is a side cross-sectional view depicting forming an etch mask over a first plurality of fin structures in a first device region of the substrate, wherein a second plurality of fin structures in a second device region of the substrate are exposed, in accordance with one embodiment of the present disclosure.

FIG. 7 is a side cross-sectional view depicting etching the second plurality of fin structures in a second device region to reduce the width of the second plurality of fin structures, in accordance with one embodiment of the present disclosure.

FIG. 8 is a side cross-sectional view depicting epitaxially forming a second semiconductor layer on the etched surface of the second plurality of fin structures in the second device region, in accordance with one embodiment of the present disclosure.

FIG. 9 is a side cross-sectional view depicting forming a gate dielectric on the first plurality of fin structures and the second plurality of fin structures, in accordance with one embodiment of the present disclosure.

FIG. 10 is a perspective view depicting one embodiment of forming gate structures and source and drain regions in direct contact with a first fin structure in the first device region and

in direct contact with a second fin structure in the second device region, in accordance with the present disclosure.

DETAILED DESCRIPTION

Detailed embodiments of the methods and structures of the present disclosure are described herein; however, it is to be understood that the disclosed embodiments are merely illustrative of the disclosed methods and structures that may be embodied in various forms. In addition, each of the examples given in connection with the various embodiments of the disclosure are intended to be illustrative, and not restrictive. Further, the figures are not necessarily to scale, some features may be exaggerated to show details of particular components. Therefore, specific structural and functional details disclosed herein are not to be interpreted as limiting, but merely as a representative basis for teaching one skilled in the art to variously employ the methods and structures of the present disclosure.

References in the specification to “one embodiment”, “an embodiment”, “an example embodiment”, etc., indicate that the embodiment described may include a particular feature, structure, or characteristic, but every embodiment may not necessarily include the particular feature, structure, or characteristic. Moreover, such phrases are not necessarily referring to the same embodiment. Further, when a particular feature, structure, or characteristic is described in connection with an embodiment, it is submitted that it is within the knowledge of one skilled in the art to affect such feature, structure, or characteristic in connection with other embodiments whether or not explicitly described. For purposes of the description hereinafter, the terms “upper”, “lower”, “right”, “left”, “vertical”, “horizontal”, “top”, “bottom”, and derivatives thereof shall relate to the invention, as it is oriented in the drawing figures. The terms “overlying”, “atop”, “positioned on” or “positioned atop” means that a first element, such as a first structure, is present on a second element, such as a second structure, wherein intervening elements, such as an interface structure, e.g. interface layer, may be present between the first element and the second element. The term “direct contact” means that a first element, such as a first structure, and a second element, such as a second structure, are connected without any intermediary conducting, insulating or semiconductor layers at the interface of the two elements.

In some embodiments, the present disclosure provides methods for forming a semiconductor device including an epitaxially deposited semiconductor layer on a low defect extremely thin semiconductor on insulator (ETSOI) layer. The term “ETSOI layer” denotes a SOI semiconductor layer having a thickness of 5 nm or less. The SOI layer may be a semiconductor layer present on the dielectric layer of a semiconductor on insulator (SOI) substrate. It has been determined that exposing an ETSOI layer to high temperatures, such as the temperatures employed to clean semiconductor materials prior to epitaxial growth thereon, can agglomerate the ETSOI layer. In some embodiments, the methods disclosed herein provide a heterostructure channel including an epitaxially grown semiconductor material on a low defect ETSOI layer by cleaning the SOI layer prior to etching to the ETSOI thickness, wherein following cleaning the SOI layer is thinned in an epitaxial deposition chamber using a halogen based etch gas. Following thinning of the SOI layer to the ETSOI thickness, the second semiconductor layer may be epitaxially grown on the etched surface. By cleaning the SOI layer with high temperature processing prior to thinning of the layer to the ETSOI thickness, the methods disclosed

herein avoid agglomeration of the ETSOI layer and provide an ETSOI layer having a low defect density.

FIGS. 1-4 depict one embodiment of a method of forming a planar semiconductor device formed on a portion of a substrate **5** that is composed of a first semiconductor layer **20a** having a thickness of less than 5 nm that is present on a buried insulating layer **15**, and a second semiconductor layer **30** having a thickness of less than 5 nm that is present on the first semiconductor layer **20a**. In some embodiments, the first semiconductor layer **20a** is composed of a semiconductor material that is different than the semiconductor material of the second semiconductor layer **30**.

As used herein, “semiconductor device” refers to an intrinsic semiconductor material that has been doped, that is, into which a doping agent has been introduced, giving it different electrical properties than the intrinsic semiconductor. Doping involves adding dopant atoms to an intrinsic semiconductor, which changes the electron and hole carrier concentrations of the intrinsic semiconductor at thermal equilibrium. Dominant carrier concentration in an extrinsic semiconductor determines the conductivity type of the semiconductor. In one embodiment, the semiconductor device is a field effect transistor.

As used herein a “field effect transistor” is a transistor in which output current, i.e., source-drain current, is controlled by the voltage applied to a gate structure. A field effect transistor has three terminals, i.e., a gate structure, source region and drain region. As used herein, the term “source” is a doped region in the semiconductor device, in which majority carriers are flowing into the channel. As used herein, the term “channel” is the region underlying the gate structure and between the source and drain of a semiconductor device that becomes conductive when the semiconductor device is turned on. As used herein, the term “drain” means a doped region in semiconductor device located at the end of the channel, in which carriers are flowing out of the transistor through the drain. A “gate structure” means a structure used to control output current (i.e., flow of carriers in the channel) of a semiconductor device through electrical or magnetic fields. The term “planar” as used to describe a semiconductor device denotes that the direction of charge carriers from the source region to the drain region of the semiconductor device is along a plane that is parallel to the upper surface of the substrate, wherein the gate structure is present on the upper surface of the substrate. Typically, in a planar device at least a portion of the source region and the drain region are present within the upper surface of the substrate.

FIG. 1 illustrates the results of the processing steps that produce a substrate **5**, i.e., semiconductor on insulator (SOI) substrate, in which the substrate **5** comprises at least a first semiconductor layer **20** (also referred to as an ETSOI layer **20**) overlying a dielectric layer **15**, wherein the first semiconductor layer **20** has a thickness of less than 10 nm. A base semiconductor layer **10** may be present underlying the dielectric layer **15**. In one embodiment, the first semiconductor layer **20** is composed of a first semiconductor material.

The first semiconductor material that provides the first semiconductor layer **20** may be any semiconducting material including, but not limited to Si, strained Si, SiC, SiGe, SiGeC, Si alloys, Ge, Ge alloys, GaAs, InAs, and InP, or any combination thereof. In one embodiment, the first semiconductor material is silicon. The first semiconductor layer **20** may be thinned to a desired thickness by planarization, grinding, wet etch, dry etch, oxidation followed by oxide etch, or any combination thereof. One method of thinning the first semiconductor layer **20** is to oxidize the Si by a thermal dry or wet oxidation process, and then wet etch the oxide layer using a

5

hydrofluoric acid mixture. This process can be repeated to achieve the desired thickness. In one embodiment, the first semiconductor layer **20** has a first thickness T1 ranging from 5.0 nm to 15.0 nm. In another embodiment, the first semiconductor layer **20** has a first thickness T1 ranging from 7.5 nm to 12.5 nm. In one example, the first semiconductor layer **20** has a first thickness T1 of 10.0 nm. The base semiconductor layer **10** may be a semiconducting material including, but not limited to Si, strained Si, SiC, SiGe, SiGeC, Si alloys, Ge, Ge alloys, GaAs, InAs, InP as well as other III/V and II/VI compound semiconductors. The base semiconductor layer **10** may have the same or a different composition than the first semiconductor layer **20**.

The dielectric layer **15** that may be present underlying the first semiconductor layer **20** and atop the base semiconductor layer **10** may be formed by implanting a high-energy dopant into a bulk semiconductor substrate and then annealing the structure to form a buried insulating layer, i.e., dielectric layer **15**. In another embodiment, the dielectric layer **15** may be deposited or grown prior to the formation of the first semiconductor layer **20**. In yet another embodiment, the substrate **5** may be formed using wafer-bonding techniques, where a bonded wafer pair is formed utilizing glue, adhesive polymer, or direct bonding.

FIG. 2 depicts one embodiment of forming a block mask **25** over a first device region **50** of the substrate **5** and etching an exposed portion of the first semiconductor layer **20a** in a second device region **55** of the substrate **5**. In one embodiment, the block mask **25** is formed protecting the portion of the substrate **5** in which the n-type semiconductor devices, e.g., nFETs, are subsequently formed. In some embodiment, the exposed portion of the substrate **5** that is not protected by the block mask **25** is subsequently processed to provide p-type semiconductor devices, e.g., pFET.

The block mask **25** may comprise soft and/or hardmask materials and can be formed using deposition, photolithography and etching. In one embodiment, the block mask **25** is a hardmask composed of a nitride-containing material, such as silicon nitride. It is noted that it is not intended that the block mask **25** be limited to only silicon nitride, as the composition of the hardmask may include any dielectric material that may be deposited by chemical vapor deposition (CVD) and related methods. Other hardmask compositions for the block mask **25** may include silicon oxides, silicon oxynitrides, silicon carbides, silicon carbonitrides, etc. Spin-on dielectrics may also be utilized as a hardmask material including, but not limited to: silsequioxanes, siloxanes, and boron phosphate silicate glass (BPSG).

In one embodiment, a block mask **25** comprising a hardmask material may be formed by blanket depositing a layer of hardmask material, providing a patterned photoresist atop the layer of hardmask material, and then etching the layer of hardmask material to provide a block mask **25** protecting at least one portion of the substrate **5**, e.g., the first device region **50** of the substrate **5**. A patterned photoresist can be produced by applying a blanket photoresist layer to the surface of the substrate **5**, exposing the photoresist layer to a pattern of radiation, and then developing the pattern into the photoresist layer utilizing resist developer. Etching of the exposed portion of the block mask **25** that is overlying the second device region **55** may include an etch chemistry for removing the exposed portion of the hardmask material and having a high selectivity to at least the block mask **25** and the first semiconductor layer **20** that is present in the second device region **55**. In one embodiment, the etch process may be an anisotropic etch process, such as reactive ion etch (RIE).

6

In some embodiments, following the formation of the block mask **25**, the exposed surface of the substrate **5**, i.e., exposed portion of the first semiconductor layer **20a**, is cleaned to remove any residual layers, foreign particles, and any residual metallic surface contamination. In one embodiment, the chemical cleaning process includes a first step of treating the exposed portion of the first semiconductor layer **20a** with hydrofluoric acid (HF), a second step of treating the surface of the exposed portion of the first semiconductor layer **20a** with a solution of ammonium hydroxide (NH₄OH) and hydrogen peroxide (H₂O₂), and a third step of treating the surface with an aqueous mixture of hydrochloric acid (HCl) and an oxidizing agent selected from the group consisting of hydrogen peroxide, ozone (O₃) and combinations thereof. The cleaning steps that include the application of the solution of ammonium hydroxide and hydrogen peroxide and the aqueous mixture of hydrochloric acid and the oxidizing agent may be provided by an RCA clean sequence.

In one embodiment and in the first step of the cleaning process, oxide material, such as silicon oxide or silicon oxynitride, is removed from the surface of the first semiconductor layer **20a** of the substrate **5** by the application of a solution of hydrofluoric acid. Hydrofluoric acid is used to etch silicon oxide (SiO₂) films on silicon substrates, because the hydrofluoric acid will etch the silicon oxide without attacking the silicon surface. The hydrofluoric acid is typically diluted with deionized (DI) water in order to slow down the etch rate of the silicon oxide, thereby ensuring better etch uniformity. In one embodiment, the dilution ratio ranges from 1:1 HF:H₂O to 300:1 H₂O:HF. In another embodiment, the hydrofluoric acid may be diluted with ammonium fluoride (NH₄F).

Following the surface treatment with hydrofluoric acid, the removal of particles and residual metallic contamination continues with an RCA clean process, which in some embodiments provides the second and third steps of the chemical cleaning process. In one embodiment, the RCA clean includes a treatment of the first semiconductor layer **20a** of the substrate **5** in a solution of ammonium hydroxide and hydrogen peroxide followed by an aqueous mixture of hydrochloric acid and an oxidizing agent (e.g., H₂O₂, O₃).

The first step of the RCA clean that includes ammonium hydroxide and hydrogen peroxide may be referred to as "SC-1" (standard clean #1). SC-1 includes a mixture of ammonium hydroxide and hydrogen peroxide and deionized water. A typical concentration ratio for the mix is 1:1:5 NH₄OH:H₂O₂:H₂O, although ratios as low as 0.05:1:5 are suitable for cleaning the substrate **5**. SC-1 typically operates in a temperature ranging from 50° C. to 70° C.

The second step of the RCA clean that includes the aqueous mixture of hydrochloric acid and an oxidizing agent may be referred to as "SC-2" (standard clean #2). SC-2 includes a mixture of hydrochloric acid, hydrogen peroxide, and deionized water. A typical concentration ratio for the mix is 1:1:5 HCl:H₂O₂:H₂O. SC-2 is typically operated in the temperature range of 50-70° C.

In another embodiment, the chemical cleaning process is provided by a hydrofluoric acid last process. In this embodiment, oxide material, such as silicon oxide or silicon oxynitride, is removed from the first semiconductor layer **20a** of the substrate **5** by the application of a solution of hydrofluoric acid. The hydrofluoric acid is typically diluted with deionized water in order to slow down the etch rate of the silicon oxide, thereby ensuring better etch uniformity. In one embodiment, the dilution ratio ranges from 1:1 HF:H₂O to 300:1 H₂O:HF. In another embodiment, the hydrofluoric acid may be diluted with ammonium fluoride (NH₄F).

The above-described chemical cleaning processes may be applied to the first semiconductor layer **20a** of the substrate **5** by methods such as immersion within a dip tank, brushing, spraying and combinations thereof. It is noted that the aforementioned methods of applying the chemical cleaning process have been provided for illustrative purposes only, and are not intended to limit the present disclosure. Any method of applying the chemical cleaning process to the first semiconductor layer **20a** of substrate **5** may be employed.

The substrate **5** may then be positioned within an epitaxial deposition chamber. The epitaxial deposition chamber includes any chamber that may be employed in epitaxial deposition. For example, the epitaxial deposition chamber may include the deposition chamber of a chemical vapor deposition (CVD) apparatus. In some embodiments, once the substrate **5** is positioned within the epitaxial deposition chamber, the first semiconductor layer **20a** of the substrate **5** may be treated with a hydrogen (H_2) containing gas at a pre-bake temperature. Typically, the hydrogen reduces the native oxide (SiO_2) containing oxygen from the surface of the first semiconductor layer **20a**. In some embodiments, treating the surface of the first semiconductor layer **20a** with a hydrogen containing gas at a pre-bake temperature provides an “oxygen-free” surface. By “oxygen-free” surface it is meant that the first semiconductor layer **20a** of the substrate **5** is free of oxide. In one embodiment, the oxygen content of the surface of the first semiconductor layer **20a** is less than 0.5%.

In one embodiment, the hydrogen containing gas is comprised of greater than 90% hydrogen (H_2). In another embodiment, the hydrogen containing gas is comprised of 100% hydrogen (H_2). It is noted that the hydrogen containing gas may include other elements, so long as the hydrogen containing gas does not provide a source of oxygen. In one embodiment, the hydrogen containing gas is applied at a pressure ranging from 10 Torr to 600 Torr.

The temperature of the hydrogen containing gas treatment is referred to as the “pre-bake temperature”. In some embodiments, in which the first semiconductor layer **20a** has not been chemically cleaned with a hydrofluoric acid last process, and there remains a native oxide, the pre-bake temperature of the hydrogen containing gas treatment may be greater than 700° C. For example, the pre-bake temperature of the hydrogen pre-bake process may be greater than 1000° C. In another example, the pre-bake temperature of the hydrogen containing gas treatment that follows the hydrofluoric acid last cleaning process may range from 1100° C. to 1200° C.

The time period for the treatment of the substrate **5** with the hydrogen containing gas may range from 15 seconds to 5 minutes. In another example, the time period for the treatment of the first semiconductor layer **20a** of the substrate **5** with the hydrogen containing gas may range from 30 seconds to 2 minutes. The above time periods are provided for illustrative purposes only and are not intended to limit the present disclosure, as other time periods may be suitable for the treatment of the first semiconductor layer **20a** of the substrate **5** with the hydrogen containing gas, so long as the time period by which the hydrogen containing gas is applied to the first semiconductor layer **20a** is sufficient to provide a substantially oxygen-free surface.

In one embodiment, the hydrogen containing gas is flown across the first semiconductor layer **20a** of the substrate **5** at a flow rate of 5000 sccm to 80000 sccm. In another embodiment, the hydrogen containing gas is flown across the first semiconductor layer **20a** of the substrate **5** at a flow rate of 24000 sccm to 35000 sccm. It is noted that the above flow

rates are provided for illustrative purposes only and may vary depending upon the configuration, e.g., size, of the epitaxial deposition chamber.

FIG. 2 further depicts etching an exposed portion of the first semiconductor layer **20a** that is present in the second device region **55** to a second thickness **T2** that is less than the first thickness **T1**, while the portion of the first semiconductor layer **20** that is present in the first device region **50** is protected by the block mask **25**. In one embodiment, the exposed portion of the first semiconductor layer **20a** is etched with a halide based gas at a temperature of less than 675° C. while the substrate **5** is present in an epitaxial deposition chamber. A “halide based gas” is a chemical compound of a halogen with a more electropositive element or group in a gaseous phase. Halogens or halogen elements are a series of nonmetal elements from Group 17 IUPAC Style (formerly: VII, VIIA) of the periodic table of elements and comprise fluorine (F), chlorine (Cl), bromine (Br), and iodine (I). Examples of halide based gasses that are suitable as an etchant for etching the exposed portion of the first semiconductor layer **20a** include chlorine gas (Cl), hydrogen fluoride (HF), hydrogen chloride (HCl), hydrogen bromide (HBr) and combinations thereof.

In one embodiment, the halide based gas that etches the first semiconductor layer **20a** in the second device region **55** of the substrate **5** further includes a carrier gas. For example, the carrier gas may be hydrogen (H_2), helium (He), argon (Ar) or nitrogen (N_2) gas. The carrier gas may comprise greater than 85% by volume of the halide based gas flow. In another embodiment, the carrier gas may comprise greater than 90% by volume of the halide based gas flow. In one embodiment, the gas flow is comprised of 90% by volume or greater of a carrier gas, such as hydrogen, 1% to 10% by volume of a halide based gas, such as hydrochloric acid (HCl).

In one embodiment, the halide based gas is applied to the exposed portion of the first semiconductor layer **20a** that is present in the second device region **55** of the substrate **5** at a flow rate ranging from 10 sccm to 20 slm. In another embodiment, the halide based gas is applied to the exposed portion of the first semiconductor layer **20a** that is present in the second device region **55** of the substrate **5** at a flow rate ranging from 100 sccm to 300 sccm. It is noted that the above flow rates are provided for illustrative purposes only and are not intended to limit the present disclosure.

The temperature of etching with the halide based gas is typically less than 675° C. In one embodiment, the etching of the exposed portion of the first semiconductor layer **20a** in the second device region **55** may range from 500° C. to 650° C. In another embodiment, the etching of the exposed portion of the first semiconductor layer **20a** in the second device region **55** may range from 550° C. to 600° C.

Following etching of the exposed portion of the first semiconductor layer **20a**, the remaining portion of the first semiconductor layer **20a** that is present in the second device region **55** has a second thickness **T2** that ranges from 1 nm to 10 nm. In another embodiment, the remaining portion of the first semiconductor layer **20a** has a second thickness **T2** that ranges from 2 nm to 5 nm.

FIG. 3 depicts epitaxially forming a second semiconductor layer **30** on the etched surface of the first semiconductor layer **20a** in the second device region **55** of the substrate **5**. The terms “epitaxial growth and/or deposition” and “epitaxially formed” mean the growth of a semiconductor material on a deposition surface of a semiconductor material, in which the semiconductor material being grown has the same crystalline characteristics as the semiconductor material of the deposition surface. In an epitaxial deposition process, the chemical

reactants provided by the source gasses are controlled and the system parameters are set so that the depositing atoms arrive at the deposition surface of the semiconductor substrate with sufficient energy to move around on the surface and orient themselves to the crystal arrangement of the atoms of the deposition surface. Thus, an epitaxial film deposited on a {100} crystal surface will take on a {100} orientation. If, on the other hand, the wafer surface has an amorphous surface layer, the depositing atoms have no surface to align to, resulting in the formation of polycrystalline or amorphous silicon instead of single crystal silicon. The temperature for epitaxial silicon deposition typically ranges from 550° C. to 900° C. Although higher temperature typically results in faster deposition, the faster deposition may result in crystal defects and film cracking.

In one embodiment, the epitaxially formed second semiconductor layer **30** may be composed of silicon-containing material. A number of different sources may be used for the deposition of epitaxial silicon. In some embodiments, the silicon containing gas sources for epitaxial growth include silane (SiH₄), disilane (Si₂H₆), trisilane (Si₃H₈), tetrasilane (Si₄H₁₀), hexachlorodisilane (Si₂Cl₆), tetrachlorosilane (SiCl₄), dichlorosilane (Cl₂SiH₂), trichlorosilane (Cl₃SiH), methylsilane ((CH₃)SiH₃), dimethylsilane ((CH₃)₂SiH₂), ethylsilane ((CH₃CH₂)SiH₃), methylidisilane ((CH₃)Si₂H₅), dimethyldisilane ((CH₃)₂Si₂H₄), hexamethyldisilane ((CH₃)₆Si₂) and combinations thereof. The temperature for epitaxial silicon deposition typically ranges from 250° C. to 900° C. Although higher temperature typically results in faster deposition, the faster deposition may result in crystal defects and film cracking.

In another embodiment, the epitaxially formed second semiconductor layer **30** may be composed of germanium (Ge). A number of different sources may be used for the deposition of epitaxial germanium. In some embodiments, the germanium containing gas sources for epitaxial growth include germane (GeH₄), digermane (Ge₂H₆), halogermane, dichlorogermane, trichlorogermane, tetrachlorogermane and combinations thereof.

In yet another embodiment, the epitaxially formed second semiconductor layer **30** is composed of a germanium-containing material, such as silicon germanium (SiGe). A number of different sources may be used for the deposition of epitaxial silicon germanium. In some embodiments, the gas source for the deposition of epitaxial SiGe may include a mixture of silicon containing gas sources and germanium containing gas sources. For example, an epitaxial layer of silicon germanium may be deposited from the combination of a silicon gas source that is selected from the group consisting of silane, disilane, trisilane, tetrasilane, hexachlorodisilane, tetrachlorosilane, dichlorosilane, trichlorosilane, methylsilane, dimethylsilane, ethylsilane, methylidisilane, dimethyldisilane, hexamethyldisilane and combinations thereof, and a germanium gas source that is selected from the group consisting of germane, digermane, halogermane, dichlorogermane, trichlorogermane, tetrachlorogermane and combinations thereof. The germanium content of the epitaxial layer of silicon germanium may range from 5% to 70%, by atomic weight %. In another embodiment, the germanium content of the epitaxial layer of silicon germanium may range from 10% to 40%.

In an even further embodiment, the epitaxially formed second semiconductor layer **30** is composed of silicon doped with carbon (Si:C). The carbon (C) content of the epitaxial grown silicon doped with carbon may range from 0.3% to 5%,

by atomic weight %. In another embodiment, the carbon content of the epitaxial grown silicon doped with carbon may range from 1% to 2%.

In one embodiment, the epitaxially formed second semiconductor layer **30** has a third thickness **T3** ranging from 1 nm to 10 nm. In another embodiment, the second semiconductor layer **30** that is present in the second device region **55** has a third thickness **T3** that ranges from 2 nm to 5 nm. In some examples, the upper surface of the epitaxially formed second semiconductor layer **30** in the second device region **55** is coplanar with the upper surface of the first semiconductor layer **20** that is present in the first device region **50**. The epitaxially formed second semiconductor layer **30** may have a defect density of less than 100 cm⁻² with as close to zero defects being preferred.

FIG. 4 depicts forming a first gate structure **60** directly on the first semiconductor layer **20** in the first device region **50** of the substrate **5**, and forming a second gate structure **65** on the second semiconductor layer **30** in the second device region **55**. Each of the first gate structure **60** and the second gate structure **65** include at least one gate dielectric **61a**, **61b** that is present on a channel portion of the substrate **5**, and at least one gate conductor **62a**, **62b** that is present on the at least one gate dielectric **61a**, **61b**. The gate conductors **62a**, **62b** may be composed of a doped semiconductor material, such as n-type doped polysilicon, or the gate conductors **62a**, **62b** may be composed of an elemental metal. When the gate conductors **62a**, **62b** are composed of an elemental metal, the gate conductors **62a**, **62b** may be any conductive metal including, but not limited to W, Ni, Ti, Mo, Ta, Cu, Pt, Ag, Au, Ru, Ir, Rh, and Re, and alloys that include at least one of the aforementioned conductive elemental metals. The gate conductors **62a**, **62b** may be composed of a single material layer or may be a multi-layered structures. The gate conductors **62a**, **62b** may be composed of the same or a different material.

The gate dielectrics **61a**, **61b** may be composed of a dielectric material, such as SiO₂, or alternatively high-k dielectrics, such as oxides of Ta, Zr, Al or combinations thereof. In another embodiment, the gate dielectrics **61a**, **61b** are comprised of an oxide, such as SiO₂, ZrO₂, Ta₂O₅ or Al₂O₃. In one embodiment, each of the gate dielectrics **61a**, **61b** has a thickness ranging from 1 nm to 10 nm. In another embodiment, each of the gate dielectric **61a**, **61b** has a thickness ranging from 1.5 nm to 2.5 nm. The gate dielectrics **61a**, **61b** may be composed of the same or a different material.

Each of the gate structures **60**, **65** may be formed using deposition, photolithography and a selective etching process. For example, the material layers that provide the gate dielectrics **61a**, **61b** and the gate conductors **62a**, **62b** may be deposited in the first device region **50** and the second device region **55** of the substrate **5**. In one embodiment, the gate dielectrics **61a**, **61b** and gate conductors **62a**, **62b** may be formed using a deposition process, such as chemical vapor deposition (CVD) or physical vapor deposition (PVD). Chemical vapor deposition (CVD) is a deposition process in which a deposited species is formed as a result of chemical reaction between gaseous reactants at greater than room temperature (25° C. to 900° C.); wherein solid product of the reaction is deposited on the surface on which a film, coating, or layer of the solid product is to be formed. Variations of CVD processes include, but not limited to, Atmospheric Pressure CVD (APCVD), Low Pressure CVD (LPCVD) and Plasma Enhanced CVD (EPCVD), Metal-Organic CVD (MOCVD) and combinations thereof may also be employed. In some embodiments, chemical vapor deposition (CVD) may be employed to deposit the dielectric material of the gate dielectrics **61a**, **61b**. In some embodiments, CVD may also be

employed to deposit the doped semiconductor material that provides the gate conductors **62a**, **62b**. In the embodiments, in which the gate conductors **62a**, **62b** are composed of a metal, the metal may be deposited using a physical vapor deposition (PVD) method, such as sputtering.

The material layers that provide the gate dielectrics **61a**, **61b**, and the gate conductors **62a**, **62b**, may then be patterned and etched to provide the first gate structure **60** and the second gate structure **65**. Specifically, a pattern is produced by applying a photoresist to the surface to be etched; exposing the photoresist to a pattern of radiation; and then developing the pattern into the photoresist utilizing a resist developer. Once the patterning of the photoresist is completed, the sections covered by the photoresist are protected while the exposed regions are removed using a selective etching process that removes the unprotected regions. Alternatively, the first and second gate structure **60**, **65** can be formed by replacement gate technology.

Still referring to FIG. 4, source and drain regions **63a**, **63b** may be formed on opposing sides of the first and second gate structures **60**, **65**. The source and drain regions **63a** that are present in the first device region **50** may be doped to an n-type conductivity when the first semiconductor layer **20** is composed of silicon, and the source and drain regions **63b** that are present in the second device region **55** may be doped to a p-type conductivity when the second semiconductor layer **30** is silicon germanium. As used herein, the term “conductivity type” denotes a dopant region being p-type or n-type. As used herein, “p-type” refers to the addition of impurities to an intrinsic semiconductor that creates deficiencies of valence electrons. In a silicon-containing substrate, examples of n-type dopants, i.e., impurities, include but are not limited to: boron, aluminum, gallium and indium. As used herein, “n-type” refers to the addition of impurities that contributes free electrons to an intrinsic semiconductor. In a silicon containing substrate examples of n-type dopants, i.e., impurities, include but are not limited to antimony, arsenic and phosphorous. The dopant that dictates the conductivity type of the source and drain regions **63a**, **63b** may be introduced to the first and second device region **50**, **55** using ion implantation. Each of the first and second device regions **50**, **55** may be selectively processes using block masks, such as a photoresist block masks. In some embodiments, prior to forming the first and second drain region **50**, **55**, a set of spacers **11** can be formed in direct contact with the sidewalls of the gate structures **60**, **65**. The spacers **11** may be employed to dictate the positioning of the dopants being implanted into the substrate **5**. In some embodiments, the first spacers **11** have a width ranging from 2.0 nm to 15.0 nm.

Referring to FIG. 4, the above process may provide a planar complementary metal oxide semiconductor (CMOS) device that includes a substrate **5** including a first device region **50** and a second device region **55**. The first device region **50** may include a first thickness T1 silicon layer, i.e., first semiconductor layer **20**, on a first portion of a buried insulating layer **15**. The second device region **55** may include a silicon germanium layer, i.e., second semiconductor **30**, on a silicon layer, i.e., etched portion of the first semiconductor layer **20a**, having a second thickness T2. The second thickness T2 of silicon layer is present on a second portion of the buried insulating layer **15**. The second thickness T2 of the silicon layer has a lesser thickness than the first thickness T1 of the silicon layer. A planar n-type semiconductor device **100**, e.g., nFET, is present in the first device region **50** of the substrate **5**, and a planar p-type semiconductor device **200**, e.g., pFET, is present in the second device region **55** of the substrate **5**.

The n-type semiconductor device **100** may be separated from the p-type semiconductor device **200** by a trench isolation region **35**.

In the embodiments, in which a p-type semiconductor device **200**, e.g., pFET, is present overlying a second semiconductor layer **30** composed of silicon germanium that is in direct contact with an etched portion of a first semiconductor layer **20a** composed of silicon, the silicon germanium material that provides the channel region of the device provides increased mobility, threshold voltage control and strain based performance enhancements. The combination of the silicon germanium second semiconductor layer **30** and the remaining portion of the silicon first semiconductor layer **20a** that is underlying the gate structure **65** may be referred to as a heterostructure channel.

FIGS. 5-10 depict one embodiment of a method of fabricating a finFET semiconductor device **300** that may include providing at least one fin structure, e.g., a first fin structure **305a**, having a first width W1 of less than 20 nm, and etching the at least one fin structure with a halide based gas at a temperature of less than 675° C. to a second width W2 that is less than the first width W1 of the at least one fin structure. The at least one fin structure may be composed of a first semiconductor material. A second semiconductor material **310** may then be epitaxially formed on an etched surface of the at least one fin structure. A gate structure **320** and source and drain regions may then be formed on the second semiconductor material **310** that is present on the at least one fin structure. In some embodiments, the at least one fin structure includes a first fin structure **305a** having the second semiconductor material **310** present thereon, and a second fin structure **305b** that is present on a second portion of the substrate that does not include the second semiconductor material. The details of this method are now discussed in greater detail.

FIG. 5 depicts an initial structure used in one embodiment of a method for forming a finFET device. The initial structure may include at least one fin structure **305a**, **305b** present atop a dielectric layer **150**. As used herein, the term “fin structure” refers to a semiconductor material, which is employed as the body of a semiconductor device, in which the gate structure is positioned around the fin structure such that charge flows down the channel on the two sidewalls of the fin structure and optionally along the top surface of the fin structure.

In one embodiment, the fin structure **305a**, **305b** and the dielectric layer **150** may be provided from an SOI substrate **90**, in which the top semiconductor layer of the SOI substrate **90** provides the fin structure **305a**, **305b**. The SOI substrate **90** typically includes a bottom semiconductor layer **101** and a top semiconductor layer **201** (also referred to as an SOI layer) that are electrically isolated from each other by a buried insulating layer (hereafter referred to as a dielectric layer **150**). The SOI layer **200** and the bottom semiconductor layer **101** may comprise at least one of Si, Ge alloys, SiGe, GaAs, InAs, InP, SiCGe, SiC as well as other III/V or II/VI compound semiconductors. The SOI layer **200** and bottom semiconductor layer **101** may comprise the same or different materials. The dielectric layer **150** separating the SOI layer **201** and the bottom semiconductor layer **101** may be a crystalline or non-crystalline oxide, nitride, oxynitride, or any other insulating material. The thickness of the various layers of the SOI substrate **90** may vary depending on the technique used in forming the same. In one embodiment, the SOI layer **201** has a thickness ranging from 3 nm to 100 nm, the dielectric layer **150** (also referred to as buried dielectric layer) has a thickness ranging from 10 nm to 150 nm, and the thickness of the bottom semiconductor layer **101** of the SOI substrate **90** may range from 10 nm to 500 nm.

In one embodiment, prior to etching the SOI substrate **90** to provide the fin structure **305a**, **305b**, a layer of the dielectric material is deposited atop the SOI substrate **90** to provide a dielectric fin cap (not shown). The material layer that provides the dielectric fin cap may be composed of a nitride, oxide, oxynitride material, and/or any other suitable dielectric layer. The dielectric fin cap may comprise a single layer of dielectric material or multiple layers of dielectric materials. The material layer that provides the dielectric fin cap can be formed by a deposition process, such as chemical vapor deposition (CVD) and/or atomic layer deposition (ALD). Alternatively, the material layer that provides the dielectric fin cap may be formed using a growth process, such as thermal oxidation or thermal nitridation. The material layer that provides the dielectric fin cap may have a thickness ranging from 1 nm to 100 nm. In one example, the dielectric fin cap is composed of an oxide, such as SiO₂, that is formed by chemical vapor deposition to a thickness ranging from 25 nm to 50 nm. In yet another embodiment, no dielectric fin cap is present.

In one embodiment, following the formation of the layer of dielectric material that provides the dielectric fin cap, a photolithography and etch process sequence is applied to the material layer for the dielectric fin cap and the SOI substrate **90**. Specifically, in one example, a photoresist mask is formed overlying the layer of the dielectric material that provides dielectric fin cap and is present overlying the SOI layer **201** of the SOI substrate **90**, in which the portion of the dielectric material that is underlying the photoresist mask provides the dielectric fin cap, and the portion of the SOI layer **201** that is underlying the photoresist mask provides the fin structure **305a**, **305b**. The exposed portions of the dielectric material that provides dielectric fin cap and the SOI layer **201**, which are not protected by the photoresist mask, are removed using a selective etch process. To provide the photoresist mask, a photoresist layer is first positioned on the layer of the dielectric material that provides dielectric fin cap. The photoresist layer may be provided by a blanket layer of photoresist material that is formed utilizing a deposition process such as, for example, CVD, PECVD, evaporation or spin-on coating. The blanket layer of photoresist material is then patterned to provide the photoresist mask utilizing a lithographic process that may include exposing the photoresist material to a pattern of radiation and developing the exposed photoresist material utilizing a resist developer.

Following the formation of the photoresist mask, an etching process may remove the unprotected portions of the dielectric material that provides the dielectric fin cap followed by removing the unprotected portion of the SOI layer **201** selective to the underlying buried insulating layer, i.e., dielectric layer **150**. For example, the transferring of the pattern provided by the photoresist into the underlying structures may include an anisotropic etch. As used herein, an "anisotropic etch process" denotes a material removal process in which the etch rate in the direction normal to the surface to be etched is greater than in the direction parallel to the surface to be etched. The anisotropic etch may include reactive-ion etching (RIE). Reactive Ion Etching (RIE) is a form of plasma etching in which during etching the surface to be etched is placed on the RF powered electrode. Moreover, during RIE the surface to be etched takes on a potential that accelerates the etching species extracted from plasma toward the surface, in which the chemical etching reaction is taking place in the direction normal to the surface. Other examples of anisotropic etching that can be used at this point of the present invention include ion beam etching, plasma etching or laser

ablation. When present, the dielectric fin cap may be removed following etching of the SOI layer **201** to form the fin structures **305a**, **305b**.

Each of the fin structures **305a**, **305b** may have a height H_1 ranging from 5 nm to 200 nm. In one embodiment, each of the fin structures **305a**, **305b** has a height H_1 ranging from 10 nm to 100 nm. In another embodiment, each of the fin structures **305a**, **305b** has a height H_1 ranging from 20 nm to 50 nm. In one embodiment, each of the fin structures **305a**, **305b** has a width W_1 of less than 20 nm. In another embodiment, each of the fin structures **305a**, **305b** has a width W_1 ranging from 3 nm to 8 nm. It is noted that any number of fin structures **305a**, **305b** may be formed.

FIG. 6 depicts forming an etch mask **315** over a first plurality of fin structures **305a** in a first device region **50** of the SOI substrate **90**, wherein a second plurality of fin structures **305b** in a second device region **55** of the SOI substrate **90** are exposed. The etch mask **315** that is depicted in FIG. 6 is similar to the etch mask **25** that is described above for protecting the first device region **50** of the substrate **5** that is depicted in FIG. 2. Therefore, the description of the etch mask **25** that is depicted in FIG. 2 is suitable for describing the etch mask **315** that is depicted in FIG. 6.

FIG. 7 depicts etching the second plurality of fin structures **305b** in a second device region **55** to reduce the width of the second plurality of fin structures **305b**. More specifically, the width of each of the second plurality of fin structures **305b** may be reduced to a second width W_2 ranging from 2 nm to 10 nm. In another embodiment, the width of each of the second plurality of fin structures **305b** may be reduced to a second width W_2 ranging from 3 nm to 6 nm. In yet another example, the width of each of the second plurality of fin structures **305b** is reduced to a second width W_2 ranging from 4 nm to 5 nm.

The etch process that reduces the width of the second plurality of fin structures **305b** may be an isotropic etch process. Contrary to an anisotropic etch process, an isotropic etch process is non-directional. Therefore, in some embodiments, the etch process that reduces the width of each of the second plurality of fin structures **305b** also reduces the height of each of the second plurality of fin structures **305b**. In some embodiments, the etch process that reduces the width of the second plurality of fin structures **305b**, reduces the height of the second plurality of fin structures **305b** to a second height H_2 of less than 10 nm. In another embodiment, the etch process that reduces the width of the second plurality of fin structures **305b**, reduces the height of the second plurality of fin structures **305b** to a second height H_2 ranging from 4 nm to 5 nm. Etching may also be done on fin structures **305a**, **305b** that have a hardmask still on the top portion of the fins, protecting them from attack. This structure is referred to as "dual gate" or "double gate" fin. The advantage is that height is not reduced during etching.

The etch process applied to the second plurality of fin structures **305b** that is depicted in FIG. 7 is similar to the etch process that is described above for reducing the thickness of the first semiconductor layer **20** that is described above with reference to FIGS. 1 and 2. Therefore, the description of the etch process that reduces the thickness of the first semiconductor layer **20** that is depicted in FIG. 2 is suitable for describing the etch process that reduces the width of the second plurality of fin structures **305b** that are depicted in FIG. 7. For example, following formation of the etch mask **315**, the exposed portion of fin structures, e.g., second plurality of fin structures **305b**, may be cleaned to remove any residual layers, foreign particles, and any residual metallic

surface contamination. The cleaning process has been described above with reference to FIG. 2.

Following the cleaning process, the SOI substrate **90** may then be positioned within an epitaxial deposition chamber. The epitaxial deposition chamber includes any chamber that may be employed in epitaxial deposition. For example, the epitaxial deposition chamber may include the deposition chamber of a chemical vapor deposition (CVD) apparatus. In some embodiments, once the SOI substrate **90** is positioned within the epitaxial deposition chamber, the second plurality of fin structures **305b** may be treated with a hydrogen (H₂) containing gas at a pre-bake temperature. Typically, the hydrogen reduces the native oxide (SiO₂) containing oxygen from the second plurality of fin structures **305b**.

In some embodiments, treating the second plurality of fin structures **305b** with a hydrogen containing gas at a pre-bake temperature provides an “oxygen-free” surface. In one embodiment, the oxygen content of the surface of the second plurality of fin structures **305b** is less than 0.5%. The details of the treatment of the second plurality of fin structures **305b** with the hydrogen-containing gas at the pre-bake temperature have been describe above with reference to FIG. 2.

Referring to FIG. 7, the second plurality of fin structures **305b** that is present in the second device region **55** are then etched to the second width **W2**, while the first plurality of fin structures **305a** that are present in the first device region **50** are protected by the block mask **315**. In one embodiment, the second plurality of fin structures **305b** are etched with a halide based gas at a temperature of less than 675° C. while the SOI substrate **90** is present in an epitaxial deposition chamber. Examples of halide based gasses that are suitable as an etchant for etching the second plurality of fin structures **305b** include chlorine gas (Cl), hydrogen fluoride (HF), hydrogen chloride (HCl), hydrogen bromide (HBr) and combinations thereof. In one embodiment, halide based gas that etches the second plurality of fin structures **305b** of the SOI substrate **90** further includes a carrier gas. For example, the carrier gas may be hydrogen (H₂), helium (He), argon (Ar) or nitrogen (N₂) gas. The carrier gas may comprise greater than 85% by volume of the halide based gas flow. In another embodiment, the carrier gas may comprise greater than 90% by volume of the halide based gas flow. In one embodiment, the gas flow is comprised of 90% by volume or greater of a carrier gas, such as hydrogen, 1% to 10% by volume of a halide based gas, such as hydrochloric acid (HCl).

In one embodiment, the halide based gas is applied to the second plurality of fin structures **305b** that are present in the second device region **55** of the substrate **5** at a flow rate ranging from 10 sccm to 20 slm. In another embodiment, the halide based gas is applied to the second plurality of fin structures **305b** at a flow rate ranging from 100 sccm to 300 sccm. It is noted that the above flow rates are provided for illustrative purposes only and are not intended to limit the present disclosure.

The temperature of etching with the halide based gas is typically less than 675° C. In one embodiment, the etching of the second plurality of fin structures **305b** may range from 500° C. to 650° C. In another embodiment, the etching of the second plurality of fin structures **305b** may range from 550° C. to 600° C.

FIG. 8 depicts epitaxially forming a second semiconductor layer **310** on the etched surface of the second plurality of fin structures **305b**. In one embodiment, the epitaxially formed second semiconductor layer **310** may be composed of silicon-containing material. A number of different sources may be used for the deposition of epitaxial silicon, such as the sources for depositing epitaxial silicon that have been described

above with reference to FIG. 3. In another embodiment, the epitaxially formed second semiconductor layer **310** may be composed of germanium (Ge). In yet another embodiment, the epitaxially formed second semiconductor layer **310** is composed of a germanium-containing material, such as silicon germanium (SiGe). The germanium content of the epitaxial layer of silicon germanium may range from 5% to 70%, by atomic weight %. In another embodiment, the germanium content of the epitaxial layer of silicon germanium may range from 10% to 40%. In an even further embodiment, the epitaxially formed second semiconductor layer **310** is composed of silicon doped with carbon (Si:C). The carbon (C) content of the epitaxial grown silicon doped with carbon may range from 0.3% to 5%, by atomic weight %. In another embodiment, the carbon content of the epitaxial grown silicon doped with carbon may range from 1% to 2%. Sources for the epitaxial deposition of germanium, silicon germanium and silicon doped with carbon (Si:C) have been described above with reference to FIG. 2.

The epitaxially formed second semiconductor layer **310** may be conformally deposited on the sidewalls and upper surfaces, i.e., etched surfaces, of the second plurality of fin structures **305b**. The term “conformal” denotes a layer having a thickness that does not deviate from greater than or less than 20% of an average value for the thickness of the layer. The term “conformally deposited” denotes a deposition method that forms a conformal layer. In one embodiment, the thickness of the epitaxially formed second semiconductor layer **310** on the sidewalls of the second plurality of fin structures **305b** is the same as the thickness of the epitaxially formed second semiconductor layer **310** on the upper surface of the second plurality of fin structures **305b**. In one embodiment, the epitaxially formed second semiconductor layer **310** has a thickness **T4** ranging from 1 nm to 10 nm. In another embodiment, the second semiconductor layer **310** that is present in the second device region **55** has a thickness **T4** that ranges from 2 nm to 5 nm. The epitaxially formed second semiconductor layer **310** may have a defect density of less than 100 cm⁻² with as close to zero defects being preferred.

Following epitaxial growth of the second semiconductor layer **310**, the block mask **315** is removed.

FIGS. 9 and 10 depict one embodiment of forming gate structures **320** and source and drain regions in direct contact with each of the plurality of first fin structures **305a** in the first device region **50** and each of the plurality of second fin structure **305b** in the second device region **55**. In one embodiment, each gate structure **320** includes a gate dielectric **321** in contact with at least the fin structure **30a**, **30b**, a gate conductor **322** on the gate dielectric **321**, and a gate dielectric cap **324** on the gate conductor **322**. The gate structure **320** may be formed by forming blanket material layers for the gate dielectric **321**, gate conductor **322**, and gate dielectric cap **324** to provide a gate stack, and patterning and etching the gate stack to provide the gate structures **320**. The gate structures **320** can be formed utilizing photolithography and etch process steps.

Referring to FIG. 9, The gate dielectrics **321** are typically positioned on at least a portion of the sidewalls of the fin structures **305a**, **305b**, but may also be formed in direct contact with the upper surface of the fin structures **305a**, **305b**. The gate dielectrics **321** may be formed by a thermal growth process such as, for example, oxidation, nitridation or oxynitridation. The gate dielectrics **321** may also be formed by a deposition process such as, for example, chemical vapor deposition (CVD), plasma-assisted CVD, metal organic chemical vapor deposition (MOCVD), atomic layer deposition (ALD), evaporation, reactive sputtering, chemical solution deposition and other like deposition processes.

The gate dielectrics **321** may be comprised of an insulating material having a dielectric constant of 4.0 or greater. In another embodiment, the gate dielectrics **321** are comprised of an insulating material having a dielectric constant greater than 7.0. The dielectric constants mentioned herein are relative to a vacuum at room temperature. In one embodiment, the gate dielectrics **321** employed in the present invention includes, but is not limited to: an oxide, nitride, oxynitride and/or silicates including metal silicates, aluminates, titanates and nitrides. In one example, when the gate dielectrics **321** are comprised of an oxide, the oxide may be selected from the group including, but not limited to SiO₂, HfO₂, ZrO₂, Al₂O₃, TiO₂, La₂O₃, SrTiO₃, LaAlO₃, Y₂O₃ and mixture thereof. The physical thickness of each of the gate dielectrics **321** may vary, but typically, the gate dielectric **321** has a thickness ranging from 1 nm to 10 nm. In another embodiment, each of the gate dielectric **321** may have a thickness ranging from 1 nm to 3 nm.

Referring to FIG. **10**, after forming the gate dielectrics **321**, and the gate conductors **322** of the gate structures **320** are formed atop the gate dielectrics **321** utilizing a deposition process, such as physical vapor deposition (PVD), CVD or evaporation. Although FIG. **10** depicts a gate structure **320** being formed on only one fin structure **305a** in the first device region **50** of the SOI substrate **90**, and depicts a gate structure **320** being formed on only one fin structure **305b** in the second device region **55** of the SOI substrate **90**, the present disclosure is not limited to only this embodiment, as any number of fin structures may be present in the first and second device regions **50**, **55** of the SOI substrate **90**.

The gate conductors **322** may be composed of polysilicon, SiGe, a silicide, a metal or a metal-silicon-nitride, such as Ta—Si—N. Examples of metals that can be used as the gate conductor **322** include, but are not limited to Al, W, Cu, Ti or other like conductive metals. In one embodiment, the gate conductor **322** comprises Ti, Zr, Hf, V, Nb, Ta, TiN, TaN or a combination thereof. The gate conductor **322** may be doped or undoped. The physical thickness of the gate conductor **322** may range from 1 nm to 10 nm. In another embodiment, the gate conductor **80** has a thickness ranging from 1 nm to 3 nm.

The gate dielectric caps **324** may be composed of a nitride, oxide, oxynitride material, and/or any other suitable dielectric layer. Gate dielectric caps **324** may comprise a single layer of dielectric material or multiple layers of dielectric materials. The dielectric layer can be formed by a deposition process, such as chemical vapor deposition (CVD) and/or atomic layer deposition (ALD). Alternatively, the gate dielectric caps **324** may be formed using a growth process, such as thermal oxidation or thermal nitridation. The gate dielectric caps **324** may have a thickness ranging from 1 nm to 100 nm. In one embodiment, the gate dielectric cap **324** is composed of a nitride, such as SiN, that is formed by chemical vapor deposition to a thickness on the order of 25 nm to 50 nm.

FIG. **10** further depicts forming a first spacer **323** in direct contact with the gate structures **320**. The first spacer **323** may be formed by depositing a conformal layer of dielectric material, such as oxides, nitrides or oxynitrides followed by etching. The first spacer **323** may have a width ranging from 1 nm to 10 nm, typically ranging from 1 nm to 5 nm.

In some embodiments, following the formation of the first spacer **323**, source and drain regions may then be formed on the exposed sidewalls of the fin structure **305a**, **305b**. The source and drain regions can be formed in the fin structures **305a**, **305b** on opposing sides of the gate structures **320** using ion implantation. In another embodiment, the source and drain regions may be formed by forming an insitu doped material on the exposed portions of the fin structures **305a**,

305b and diffusing dopant from insitu doped material into the sidewalls of the fin structures **305a**, **305b**. In one embodiment, the source and drain regions of the fin structures **305a** that are present in the first device region **50** may be doped to an n-type conductivity when the first semiconductor material of the fin structure **305a** in the first device region **50** is composed of silicon. In this embodiment, the finFET device that is formed in the first device region **50** is an n-type finFET **400**. In one embodiment, the source and drain regions that are present in the second device region **55** may be doped to a p-type conductivity when the second semiconductor layer **310** on the second fin structure **305b** is silicon germanium. In this embodiment, the finFET device that is formed in the second device region **55** is a p-type finFET **300**. Each of the first and second device regions **50**, **55** may be selectively processed using block masks, such as a photoresist block masks.

In the embodiments, in which a p-type finFET device **300** is formed on a second semiconductor layer **310** of silicon germanium that is in direct contact with an etched portion of a fin structure **305b** composed of silicon, the silicon germanium material that provides the channel region of the device provides increased mobility, threshold voltage control and strain based performance enhancements.

While the present disclosure has been particularly shown and described with respect to preferred embodiments thereof, it will be understood by those skilled in the art that the foregoing and other changes in forms and details may be made without departing from the spirit and scope of the present invention. It is therefore intended that the present invention not be limited to the exact forms and details described and illustrated, but fall within the scope of the appended claims.

What is claimed is:

1. A method of fabricating a semiconductor device comprising:

providing a substrate comprising at least a first semiconductor layer atop an entire topmost surface of a dielectric layer, wherein the first semiconductor layer has a first thickness of less than 10 nm;

etching the first semiconductor layer with a halide based gas to a second thickness that is less than the first thickness;

epitaxially forming a second semiconductor layer directly on a topmost surface of an etched portion of the first semiconductor layer, wherein the entire thickness of the second semiconductor layer is less than said first thickness of the first semiconductor layer;

forming a gate structure directly on a topmost surface of the second semiconductor layer; and
forming a source region and a drain region on opposing sides of the gate structure.

2. The method of claim **1**, wherein prior to etching the first semiconductor layer is treated with a hydrogen containing gas at a pre-bake temperature that is greater than 700° C.

3. The method of claim **1**, wherein the etching of the first semiconductor layer with the halide based gas comprises a hydrogen chloride (HCl) gas, a chlorine (Cl) gas or a combination thereof.

4. The method of claim **3**, wherein the hydrogen chloride (HCl) gas is applied to the semiconductor layer at a flow rate ranging from 10 sccm to 20 slm, or the chlorine (Cl) gas is applied to the semiconductor layer at a flow rate ranging from 10 sccm to 300 sccm.

5. The method of claim **1**, wherein the second thickness of the first semiconductor layer ranges from 1.0 nm to 5.0 nm.

6. The method of claim 1, wherein the first semiconductor layer is silicon and the second semiconductor layer is a germanium-containing semiconductor.

7. The method of claim 1, wherein the second semiconductor layer has a thickness ranging from 2 nm to 5 nm. 5

8. The method of claim 1, wherein the semiconductor device is a complementary metal oxide semiconductor (CMOS) device and the substrate comprises a first device region and a second device region separated by an isolation region. 10

9. The method of claim 8, wherein an etch mask is formed over the first device region, wherein during the etching of the first semiconductor layer the etch mask protects the first semiconductor layer in the first device region with the first semiconductor layer in the second device region is etched to the second thickness, and the etch mask is present over the first device region during the epitaxially forming of the second semiconductor layer on the etched surface of the first semiconductor layer. 15

10. The method of claim 9, wherein the first semiconductor layer is silicon and the second semiconductor layer is silicon germanium, wherein following the epitaxially forming of the second semiconductor layer in the second device region of the substrate, the etch mask is removed and a n-type field effect transistor is formed on the first semiconductor layer in the first device region of the substrate. 20 25

11. The method of claim 1, wherein said etching the first semiconductor layer occurs at a temperature of between 500° C. to 650° C., inclusive. 30

* * * * *